1980:41660 CAPLUS AN

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Peptidic alkaloids. X. Approach for the synthesis of peptidic alkaloids. 1. Reactivity of N-tolylsulfonylaziridines towards reactive nucleophiles

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AB In order to realize the stereospecific introduction of a phenol ether function at the β-position of amino acids, present in peptidic alkaloids, the regiochem. of ring opening reactions of the aziridines I (R = H, Me, CHMe2, Ph), obtained from the corresponding RCH(OH)CH/(NH2)CO2H by a reported procedure (Okawa, K., et al., 1972), within nucleophiles (Man3, p-MeC6H4ONa, NH3, p-MeC6H4OH) was examd. Only I (R = Ph) was cleaved regioselectively in each case to give p-MeC6H4NHCHPhCHRCONHPh (R = N3, OC6H4Me-p). The results are discussed in terms of electronic and steric factors.

IT 70275-44-4P 70275-50-2P 70275-53-5P

RL: SPN (Synthetic preparation); PREP (Preparation)
(prepn. of)

RN 70275-44-4 CAPLUS

CN Benzenepropanamide, β -azido- α -[[(4-methylphenyl)sulfonyl]amino]-N-phenyl-, (R*,R*)- (9CI) (CA INDEX NAME)

Relative stereochemistry.

RN 70275-50-2 CAPLUS

CN Benzenepropanamide, α -azido- β -[[(4-methylphenyl)sulfonyl]amino]-N-phenyl-, (R*,R*)- (9CI) (CA INDEX NAME)

Relative stereochemistry.

RN 70275-53-5 CAPLUS

CN Benzenepropanamide, α -(4-methylphenoxy)- β -[[(4-methylphenyl)sulfonyl]amino]-N-phenyl-, (R*,R*)- (9CI) (CA INDEX NAME)

Relative stereochemistry.